

GenCore version 4.5
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OM protein - protein search, using sw model
Run on: October 1, 2002, 06:24:15 ; Search time 32.44 seconds
(without alignments)
1222.360 Million cell updates/sec

Title: US-09-522-752-2
Perfect score: 1854
Sequence: 1 MADDYGESESTSSMEDYVNFN.....EGSLKLSSMLLETSGALS 357

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 747574 seqs, 111073796 residues

Tc number of hits satisfying chosen parameters: 747574

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_032802.*
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22: /SIDS1/gcgdata/hold-geneseg/geneseq-emb1/AA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES									
Result No.	Score	Query Match	Length	ID	Description				
1	1854	100.0	357	21	AA19605	Human CC chemokine			
2	1854	100.0	357	21	AA19605	Human G protein-co			
3	1854	100.0	357	22	AA19605	Human CCR9b protei			
4	1854	100.0	369	22	AA19605	Human CCR9a protei			
5	1848	99.7	357	22	AA19605	Human mutant G pro			
6	1848	99.7	369	22	AA19605	Non-endogenous hum			
7	780.5	42.1	358	15	AA19605	Partial sequence o			
8	780.5	42.1	358	21	AA19605	Human 7TM recepto			
9	780.5	42.1	378	19	AA19605	Human V31 seven tr			
10	780.5	42.1	378	21	AA19605	Human 7TM recepto			
11	780.5	42.1	378	21	AA19605	Human G protein-co			

12	780.5	42.1	378	22	AA19605	Human CCR7 protein
13	780.5	42.1	378	22	AA19605	Human CCR7. Homo
14	780.5	42.1	410	15	AA19605	Putative seven tra
15	780.5	42.1	410	15	AA19605	Polypeptide sequen
16	780.5	42.1	410	21	AA19605	Genomic clone of 7
17	780.5	42.1	569	22	AA19605	Novel human diagno
18	777.5	41.9	378	15	AA19605	Putative seven tra
19	776.5	41.9	378	21	AA19605	Human mutant G pro
20	758.5	40.9	378	15	AA19605	Epstein Barr virus
21	758.5	40.9	378	19	AA19605	G-protein coupled
22	758.5	40.9	378	19	AA19605	Epstein Barr virus
23	757.5	40.9	378	21	AA19605	7TM receptor prote
24	742	40.0	359	15	AA19605	Seven transmembran
25	742	40.0	359	19	AA19605	Murine V31 seven t
26	742	40.0	359	21	AA19605	Murine 7TM recepto
27	721	38.9	361	20	AA19605	An Epstein-barr vi
28	671.5	36.2	351	20	AA19605	A7 times membrane
29	652	35.2	350	22	AA19605	Amino acid sequenc
30	640	34.5	350	20	AA19605	Mouse BGCKr protei
31	638.5	34.4	369	22	AA19605	Human CCR6b protei
32	638.5	34.4	374	22	AA19605	Human CCR6a protei
33	637	34.4	349	20	AA19605	Human HFIA041 prot
34	637	34.4	350	20	AA19605	Human BGCKr protei
35	637	34.4	350	20	AA19605	A human seven-pass
36	637	34.4	350	20	AA19605	Human signal pepti
37	637	34.4	350	20	AA19605	Human HFIA041 prot
38	637	34.4	350	21	AA19605	Human seven transm
39	637	34.4	350	22	AA19605	Human CCR11 protei
40	637	34.4	350	22	AA19605	Human G protein-co
41	637	34.4	382	22	AA19605	Amino acid sequenc
42	637	34.4	382	22	AA19605	Human chemokine re
43	635	34.3	333	20	AA19605	Human BGCKr, partia
44	632	34.1	350	21	AA19605	Human orphan G pro
45	632	34.1	350	21	AA19605	Human G protein co

ALIGNMENTS

RESULT 1	
AA19605	
ID	AA19605 standard; Protein; 357 AA.
XX	
AC	AA19605;
XX	
DT	22-JAN-2001 (first entry)
XX	
DE	Human CC chemokine receptor GPR-9-6.
XX	
KW	GPR-9-6; human; chemokine receptor; TECK; cancer; leukaemia;
KW	lymphoma; carcinoma; inflammation; Crohn's disease; colitis;
KW	therapy; diagnosis.
XX	
OS	Homo sapiens.
XX	
FN	WO200053635-A1.
XX	
PD	14-SEP-2000.
XX	
PF	10-MAR-2000; 2000WO-US06240.
XX	
PR	11-MAR-1999; 99US-0266464.
XX	
PA	(LEUK-) LEUKOSITE INC.
XX	
PI	Andrew DP, Zabel BA, Ponath PD;
XX	
DR	WPI; 2000-572263/53.
XX	
PT	Antibody or its antigen-binding fragment which binds to the mammalian
PT	CC chemokine receptor GPR-9-6, useful for treating inflammatory
PT	diseases, cancer or inhibiting GPR-9-6-mediated homing of leukocytes to
PT	mucosal tissue -

XX PS Disclosure; Fig 14A-B; 114pp; English.

XX CC The present sequence is that of human GPR-9-6, a CC chemokine

CC CC receptor that is expressed on the majority of thymocytes and also

CC CC on a subset of memory CD4 lymphocytes that traffic to mucosal

CC CC sites, suggesting a dual role in T cell development and mucosal

CC CC immune response. The invention relates to an antibody that binds

CC CC to GPR-9-6 and blocks the binding of a ligand, such as TECK (see

CC CC AAB19607), to the receptor. Also provided is a method of identifying

CC CC agents which can bind to GPR-9-6 and inhibit the binding of a

CC CC ligand and/or modulate a function of GPR-9-6. The antibodies can

CC CC be used to detect or measure expression of GPR-9-6 receptor. They

CC CC are useful for treating an inflammatory disease, cancer and

CC CC inhibiting GPR-9-6-mediated homing of leukocytes to mucosal tissue.

CC CC The cancer treated is acute or chronic leukaemia (e.g., acute T-cell

CC CC lymphoblastic leukaemia, acute B-cell lymphoblastic leukaemia,

CC CC chronic T-cell lymphoblastic leukaemia, chronic B-cell lymphoblastic

CC CC leukaemia), lymphoma (e.g., Hodgkin's disease, T cell lymphoma) or

CC CC carcinoma (e.g. breast, melanoma, myeloma, or adenoma). The

CC CC inflammatory diseases treated are Crohn's disease, colitis

CC CC (claimed), inflammatory bowel disease, mastitis, vaginitis,

CC CC cholangitis or pericholangitis, chronic bronchitis, asthma, graft

CC CC versus host disease, hypersensitivity pneumonitis, collagen

CC CC diseases, sarcoidosis, and other idiopathic conditions. Other

CC CC diseases that can be treated by the antibodies are autoimmune

CC CC diseases (e.g. rheumatoid arthritis, multiple sclerosis), infectious

CC CC diseases (e.g. bacterial and viral infections), atherosclerosis,

CC CC restenosis, AIDS, pancreatitis, insulin-dependent diabetes mellitus,

CC CC and diseases in which angiogenesis or neovascularization play a role.

XX CC Sequence 357 AA;

XX CC

Query Match 100.0%; Score 1854; DB 21; Length 357;

Best Local Similarity 100.0%; Pred. No. 1.4e-191;

Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MADDYGSSTSSMEDYVNFDFYCEKNNVRQFASHFLPPLYWLVTIVGALNSLVILV 60

Db 1 maddygsstssmedyvfnfdyfceknvrvqfashflpplwlvfivgalnslvllv 60

Qy 61 YWYCTRVKTMDFMLNLAIADLLFLVTLFPWATAAADQWKFQFMCKVNVSMYKMFYS 120

Db 61 ywycrvktmdtmfmlnlaiadllflvtpfwataaadqwkfqfcmckvvnsmymkmfys 120

Qy 121 CVLLIMCISVDRIYAIQAAMRAHTWREKRLYSKMVCFIIVWLAALCIPILYSQIKEE 180

Db 121 cvllimcisdvryiaiaqamrahtwrekrllyskmvctfiwvlaaalcipeillysqikee 180

Qy 181 SGIAICTMVYPSDESTKLKSAVLTLKVLGFFLPFVVMACCYTIITHTLIOAKSSKHKA 240

Db 181 sgiaictmvpsdestklksavtlklvlgfflpfvvmaccytiithtliqaksskhka 240

Qy 241 LKVTITVTLTVFLVQFPYNCILLVQTIDAYAMFISNCAVSTNIDICQVOTIAFFHSCL 300

Db 241 lkvtitvltvflvqfpyncillvgtidayamfiscavstnidicqvtiaffhscl 300

Qy 301 NPVLVYVGERFRDLVKTLKLCISQAQWVSFTTRREGSKLSSMLLETTSGLSL 357

Db 301 npvlvyvgerfrdlvktlknlgcisqaqwwsfttrregsklssmllettsgalsl 357

RESULT 2

AAAY90615

ID AAAY90615 standard; Protein; 357 AA.

XX CC

AC AAAY90615;

XX CC

DT 21-AUG-2000 (first entry)

XX CC

DE Human G protein-coupled receptor GPR9-6.

XX CC

KW G protein-coupled receptor; GPCR; constitutively active; intracellular loop 3; transmembrane domain 6; drug screening; agonist; antagonist.

XX Homo sapiens.

OS WO200022129-A1.

PN 20-APR-2000.

XX 12-OCT-1999; 99WO-US23938.

PR 13-OCT-1998; 98US-0170496.

XX (AREN-) ARENA PHARM INC.

XX Behan DP, Chalmers DT, Liaw CW;

DR WPT; 2000-329165/28.

DR N-PSDB; AAA30596.

XX Non-endogenous constitutively activated human G protein-coupled receptors, useful for identifying agonists for use as pharmaceutical agents

PT agents

XX Example 1; Page 119-120; 341pp; English.

XX The invention relates to constitutively active, non-endogenous versions of endogenous human orphan G protein-coupled receptors (GPCRs, AAAY90643-AAAY90677 and AAAY90683-Y90687), and to DNA encoding them (AAAY30709-A30743 and AAAY30775-A30779). The mutant proteins of the invention contain a mutation in a portion of the protein comprising intracellular loop 3 (IC3) and transmembrane domain 6 (TM6). A non-endogenous amino acid, X, is substituted for an endogenous residue in IC3 at a position 16 amino acids N-terminal of an endogenous residue in TM6 to form a sequence X-(AA)15-Pro. The endogenous amino acid is selected from Lys, His, Arg or Ala, and is preferably Lys. When the endogenous residue at this position is Lys, this residue is replaced by His, Arg or preferably Ala. The 15 amino acid stretch between the substituted amino acid and the Pro may be endogenous, non-endogenous, or a mixture of endogenous and non-endogenous residues. The constitutively active GPCRs are useful for identifying antagonists, agonists and partial agonists for use as pharmaceutical agents. The mutant proteins are also useful in research settings for elucidating the roles of the receptors in normal and diseased conditions. Antagonists for a particular GPCR are useful for treating diseases and disorders associated with that receptor. Because the novel mutant GPCRs are constitutively active, they can be used directly for screening of compounds without the need for endogenous ligands. The present sequence represents a human wild-type GPCR referred to in an exemplification of the invention.

XX Sequence 357 AA;

XX

Query Match 100.0%; Score 1854; DB 21; Length 357;

Best Local Similarity 100.0%; Pred. No. 1.4e-191;

Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MADDYGSSTSSMEDYVNFDFYCEKNNVRQFASHFLPPLYWLVTIVGALNSLVILV 60

Db 1 maddygsstssmedyvfnfdyfceknvrvqfashflpplwlvfivgalnslvllv 60

Qy 61 YWYCTRVKTMDFMLNLAIADLLFLVTLFPWATAAADQWKFQFMCKVNVSMYKMFYS 120

Db 61 ywycrvktmdtmfmlnlaiadllflvtpfwataaadqwkfqfcmckvvnsmymkmfys 120

Qy 121 CVLLIMCISVDRIYAIQAAMRAHTWREKRLYSKMVCFIIVWLAALCIPILYSQIKEE 180

Db 121 cvllimcisdvryiaiaqamrahtwrekrllyskmvctfiwvlaaalcipeillysqikee 180

Qy 181 SGIAICTMVYPSDESTKLKSAVLTLKVLGFFLPFVVMACCYTIITHTLIOAKSSKHKA 240

Db 181 sgiaictmvpsdestklksavtlklvlgfflpfvvmaccytiithtliqaksskhka 240

QY 241 LKVTITVTLTVFVLSQFPYNCILLVQTIDAYAMFISNCAVSTNIDICQVOTOTIAFFHSL 300
 Db 241 lkvtitvltvfvlsqfpyncillvgtidayamfiscavstnidicfqvtqti affhsl 300
 QY 301 NPVLYVFGFRFRDLVTKLNLGICISQAQWVSTRREGSKLSMLETTSGLSL 357
 Db 301 npvlyvfgfrfrdlvktlknlgisqagwvstrregsklsmlettsgalsl 357

RESULT 3

AAG80117
 ID AAG80117 standard; Protein; 357 AA.
 XX
 AC AAG80117;
 XX
 DT 17-JAN-2002 (first entry)
 DE Human CCR9b protein.
 XX
 KW Chemokine; tumour diagnosis; colorectal; prostatic; organ rejection;
 KW inflammation; autoimmune disease; metastasis; bronchial asthma; lupus;
 KW chronic bowel inflammation; rheumatoid arthritis; cytostatic;
 KW antiinflammatory; antiasthmatic; immunosuppressive; dermatological;
 KW antirheumatic; antiarthritic.
 XX
 OS Homo sapiens.
 XX
 PN WO200172830-A2.
 XX
 PD 04-OCT-2001.
 XX
 PF 02-APR-2001; 2001WO-EP03708.
 XX
 PR 31-MAR-2000; 2000DE-1016013.
 XX
 PA (IPFP-) IPF PHARM GMBH.
 PA (FORS/) FORSMANN U.
 XX
 PI Forssmann W, Adermann K, Heitland A, Spodsborg N;
 XX
 DR WPI; 2001-626256/72.
 XX
 PT Diagnostic agent containing two or more receptor-specific ligands,
 PT useful for detecting tumors, inflammation etc., also therapeutic use of
 PT ligand inhibitors -
 XX
 PS Disclosure; Page 11; 26pp; German.

XX This invention describes a novel diagnostic agent (A) comprising at least
 CC two different ligands (I) for receptors (II) that are implicated in
 CC disease. (A) are used for the diagnosis of tumors (especially colorectal
 CC or prostatic), organ rejection, inflammation and autoimmune diseases.
 CC Also inhibitors of (I) are used therapeutically against tumors (and their
 CC metastases), inflammation (particularly bronchial asthma or chronic bowel
 CC inflammation), or autoimmune diseases (rheumatoid arthritis or lupus),
 CC where the (cardio)vascular, lymphatic, respiratory, nervous, digestive,
 CC endocrine, motor or urogenital systems or skin are affected, and bone
 CC marrow diseases. The products of the invention are chemokine derivatives
 CC which have cytostatic, antiinflammatory, antiasthmatic,
 CC immunosuppressive, dermatological, antirheumatic, antiarthritic.
 CC Chemokines act on specific tumor and inflammatory cells through a
 CC constellation of chemokine receptors (CR), which control migration and
 CC proliferation of these cells. AAG80045-AAG80128 represent human chemokine
 CC fragments used to illustrate the method of the invention.

Sequence 357 AA;

Query Match 100.0%; Score 1854; DB 22; Length 357;
 Best Local Similarity 100.0%; Pred. No. 1.4e-191;
 Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MADDYGSESTSSMEDYVNFNFTDFYCEKNVVRQFASHFLPPLYLWLVFIVGALGNSLIVLY 60
 Db 1 maddygsestssmedyvnfnftdfyceknvvrqfashflpplylwlvfivgalgnslylv 60
 QY 61 YWYCTRVKVTMDMFLNLALADLLFLVLPFWAIAAADQWKFQFMCKVNVSMYKMFYS 120
 Db 61 ywyctrvktmdmflnlaladllflvlpfwaiaaadqwkfqfmckvvnsmymkmfys 120
 QY 121 CVLLIMCISVDTRYIAIAQAMRAHTWRKRLLYSKWVCFTHWLAALCIPILYSQIKEE 180
 Db 121 cvllimcislvdryiaiaqamrahtwrekrillyskwvcftiwlalaaicpellysqikee 180
 QY 181 SGIAICTWVYPSDESTKLKSAVLTLKVLGFLPFPVVMACCYTHIHTLOAKKSSKHKA 240
 Db 181 sgiaicmwypsdetklksavltklvlgfplfpvvmaccythihltloakkskhka 240
 QY 241 LKVTITVTLTVFVLSQFPYNCILLVQTIDAYAMFISNCAVSTNIDICQVOTOTIAFFHSL 300
 Db 241 lkvtitvltvfvlsqfpyncillvgtidayamfiscavstnidicfqvtqti affhsl 300
 QY 301 NPVLYVFGFRFRDLVTKLNLGICISQAQWVSTRREGSKLSMLETTSGLSL 357
 Db 301 npvlyvfgfrfrdlvktlknlgisqagwvstrregsklsmlettsgalsl 357
 RESULT 4
 AAG80116
 ID AAG80116 standard; Protein; 369 AA.
 XX
 AC AAG80116;
 XX
 DT 17-JAN-2002 (first entry)
 DE Human CCR9a protein.
 XX
 KW Chemokine; tumour diagnosis; colorectal; prostatic; organ rejection;
 KW inflammation; autoimmune disease; metastasis; bronchial asthma; lupus;
 KW chronic bowel inflammation; rheumatoid arthritis; cytostatic;
 KW antiinflammatory; antiasthmatic; immunosuppressive; dermatological;
 KW antirheumatic; antiarthritic.
 XX
 OS Homo sapiens.
 XX
 PN WO200172830-A2.
 XX
 PD 04-OCT-2001.
 XX
 PF 02-APR-2001; 2001WO-EP03708.
 XX
 PR 31-MAR-2000; 2000DE-1016013.
 XX
 PA (IPFP-) IPF PHARM GMBH.
 PA (FORS/) FORSMANN U.
 XX
 PI Forssmann W, Adermann K, Heitland A, Spodsborg N;
 XX
 DR WPI; 2001-626256/72.
 XX
 PT Diagnostic agent containing two or more receptor-specific ligands,
 PT useful for detecting tumors, inflammation etc., also therapeutic use of
 PT ligand inhibitors -
 XX
 PS Disclosure; Page 11; 26pp; German.
 XX
 CC This invention describes a novel diagnostic agent (A) comprising at least
 CC two different ligands (I) for receptors (II) that are implicated in
 CC disease. (A) are used for the diagnosis of tumors (especially colorectal
 CC or prostatic), organ rejection, inflammation and autoimmune diseases.
 CC Also inhibitors of (I) are used therapeutically against tumors (and their
 CC metastases), inflammation (particularly bronchial asthma or chronic bowel
 CC inflammation), or autoimmune diseases (rheumatoid arthritis or lupus),
 CC where the (cardio)vascular, lymphatic, respiratory, nervous, digestive,
 CC endocrine, motor or urogenital systems or skin are affected, and bone

CC marrow diseases. The products of the invention are chemokine derivatives
 CC which have cytostatic, antiinflammatory, antitumorigenic, antithrombotic,
 CC immunosuppressive, dermatological, antirheumatic, antiarthritic.
 CC Chemokines act on specific tumor and inflammatory cells through a
 CC constellation of chemokine receptors (CR), which control migration and
 CC proliferation of these cells. AAG80045-AAG80128 represent human chemokine
 CC fragments used to illustrate the method of the invention.
 XX
 SQ Sequence 369 AA;

Query Match 100.0%; Score 1854; DB 22; Length 369;
 Best Local Similarity 100.0%; Pred. No. 1.4e-191;
 Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MADDYGGSESTSMEDYVNFNFTDFYCEKNVQFASHFLPPLYWLVFVIGALNSLVILV 60
 DB 13 maddygsestsmedyvnnftdfyceknvqfashflpplwlvfivgalslvilv 72
 QY 61 YWYCTRVKTMDFLLNLAIADLLFLVLPFWAIAAADQWKFQTFMCKVNSMYKMFYS 120
 DB 73 ywycrvktmdmflnlaiadllflvlpfwaiaaqqkftfcmckvnsmykmfys 132
 QY 121 CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKMVCFTIWWLAALCIPPEILYSQIKEE 180
 DB 133 cvllimcisdryiaiaqamrahtwrekrllyskmvctiwwlaaalcipeiylsqikee 192
 QY 181 SGIAICTMVYPSDESTKLKSAVLTLLKVLGFPFVWVACCVTIIHTLIQAKSSKHKA 240
 DB 193 sgiaictmvpsdestklksavltllkvlgfflvfvmvaccytlilhtliqaksskhka 252
 QY 241 LKVTITVTLVFLVSQFPYNCILLVQTDAYAMFISNCVSTNIDICFQVTTOTIAFFHSCL 300
 DB 253 lkvtitvltvflvsqfpyncillvqtdayamfiscavstnidicfvttotiaffhscl 312
 QY 301 NPVLVYVGFGRFRDLVKTLLKGLCISQAQWVSFTRREGSLKLSMLLETTSGALS 357
 DB 313 npvlvyvgerfrdvlktllkglcisaqawvsftrregsklslmleettsgalsl 369

RESULT 5
 AAY90649
 ID AAY90649 standard; Protein: 357 AA.

XX
 AC AAY90649;
 XX
 DT 21-AUG-2000 (first entry)
 XX
 TX Human mutant G protein-coupled receptor GPR9-6 (L241K).
 KW G protein-coupled receptor; GPCR; constitutively active;
 KW intracellular loop 3; transmembrane domain 6; drug screening;
 KW agonist; antagonist; mutant; mutein.

XX Homo sapiens.
 OS Synthetic.

XX WO200022129-A1.
 XX
 PD 20-APR-2000.

XX 12-OCT-1999; 99WO-US23938.
 XX
 PR 13-OCT-1998; 98US-0170496.
 XX
 PA (AREN-) ARENA PHARM INC.

PI Behan DP, Chalmers DT, Liaw CW;
 XX
 DR WPI; 2000-329165/28.
 DR N-PSDB; AAA30715.

XX Non-endogenous constitutively activated human G protein-coupled
 PT

PT receptors, useful for identifying agonists for use as pharmaceutical
 PT agents
 XX
 PS Example 2; Page 226-227; 34lpp; English.

XX The invention relates to constitutively active, non-endogenous versions
 CC of endogenous human orphan G protein-coupled receptors (GPCRs, AAY90643-
 CC AAY90677 and AAY90683-Y90687), and to DNA encoding them (AAA30709-A30743
 CC and AAA30775-A30779). The mutant proteins of the invention contain a
 CC mutation in a portion of the protein comprising intracellular loop 3
 CC (IC3) and transmembrane domain 6 (TM6). A non-endogenous amino acid, X,
 CC is substituted for an endogenous residue in IC3 at a position 16 amino
 CC acids N-terminal of an endogenous proline in TM6 to form a sequence
 CC X-(AA)15-Pro. The endogenous amino acid is selected from Lys, Arg,
 CC or Ala, and is preferably Lys. When the endogenous residue at this
 CC position is Lys, this residue is replaced by His, Arg or preferably Ala.
 CC The 15 amino acid stretch between the substituted amino acid and the pro
 CC may be endogenous, non-endogenous, or a mixture of endogenous and
 CC non-endogenous residues. The constitutively active GPCRs are useful for
 CC identifying antagonists, agonists and partial agonists for use as
 CC pharmaceutical agents. The mutant proteins are also useful in research
 CC settings for elucidating the roles of the receptors in normal and
 CC diseased conditions. Antagonists for a particular GPCR are useful for
 CC treating diseases and disorders associated with that receptor. Because
 CC the novel mutant GPCRs are constitutively active, they can be used
 CC directly for screening of compounds without the need for endogenous
 CC ligands. Sequences AAY90643- AAY90677 and AAY90683-Y90687 the mutant
 CC human GPCRs of the invention.

XX Sequence 357 AA;

Query Match 99.7%; Score 1848; DB 21; Length 357;
 Best Local Similarity 99.7%; Pred. No. 6.2e-191;
 Matches 356; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MADDYGGSESTSMEDYVNFNFTDFYCEKNVQFASHFLPPLYWLVFVIGALNSLVILV 60
 DB 1 maddygsestsmedyvnnftdfyceknvqfashflpplwlvfivgalslvilv 60

QY 61 YWYCTRVKTMDFLLNLAIADLLFLVLPFWAIAAADQWKFQTFMCKVNSMYKMFYS 120
 DB 61 ywycrvktmdmflnlaiadllflvlpfwaiaaqqkftfcmckvnsmykmfys 120

QY 121 CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKMVCFTIWWLAALCIPPEILYSQIKEE 180
 DB 121 cvllimcisdryiaiaqamrahtwrekrllyskmvctiwwlaaalcipeiylsqikee 180

QY 181 SGIAICTMVYPSDESTKLKSAVLTLLKVLGFPFVWVACCVTIIHTLIQAKSSKHKA 240
 DB 181 sgiaictmvpsdestklksavltllkvlgfflvfvmvaccytlilhtliqaksskhka 240

QY 241 LKVTITVTLVFLVSQFPYNCILLVQTDAYAMFISNCVSTNIDICFQVTTOTIAFFHSCL 300
 DB 241 lkvtitvltvflvsqfpyncillvqtdayamfiscavstnidicfvttotiaffhscl 300

QY 301 NPVLVYVGFGRFRDLVKTLLKGLCISQAQWVSFTRREGSLKLSMLLETTSGALS 357
 DB 301 npvlvyvgerfrdvlktllkglcisaqawvsftrregsklslmleettsgalsl 357

RESULT 6
 ABB56344
 ID ABB56344 standard; Protein: 369 AA.

XX
 AC ABB56344;
 XX

DT 18-FEB-2002 (first entry)

XX Non-endogenous human GPCR protein, SEQ ID NO: 481.

XX Human; G protein-coupled receptor; GPCR; non-endogenous; mutant;
 KW constitutively activated GPCR; agonist; disease.

CC acids N-terminal of an endogenous proline in TM6 to form a sequence
CC X-(AA)15-Pro. The endogenous amino acid is selected from Lys, His, Arg
CC or Ala, and is preferably Lys. When the endogenous residue at this
CC position is Lys, this residue is replaced by His, Arg or preferably Ala.
CC The 15 amino acid stretch between the substituted amino acid and the Pro
CC may be endogenous, non-endogenous, or a mixture of endogenous and
CC non-endogenous residues. The constitutively active GPCRs are useful for
CC identifying antagonists, agonists and partial agonists for use as
CC pharmaceutical agents. The mutant proteins are also useful in research
CC settings for elucidating the roles of the receptors in normal and
CC diseased conditions. Antagonists for a particular GPCR are useful for
CC treating diseases and disorders associated with that receptor. Because
CC the novel mutant GPCRs are constitutively active, they can be used
CC directly for screening of compounds without the need for endogenous
CC ligands. The present sequence represents a human wild-type GPCR referred
CC to in an exemplification of the invention.

XX Sequence 378 AA;

Query Match 42.1%; Score 780.5; DB 21; Length 378;
Best Local Similarity 42.9%; Pred. No. 1.2e-75;
Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

QY 1 MADDYGESESSMEDYVNFDFCYCKNNVRQFASHFLPPLYWLVFIVGALGNSLVILV 60
DB 28 vtdydgntt-----vdytlfeslcskdkvrnfkawlplmysislfcvlgilnglvlt 82
QY 61 YWYTRVKTMTDFLLNLAIADLLFLVLPFWATAAADQMKFQFMCKVNSYKMFYS 120
DB 83 yiyfkrktmtdtyllnlavadiilflitlpfwaysaakswvfgvhfcklifaikmsffs 142
QY 121 CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKMVCFTIWLAAALCIPILYSQIKEE 180
DB 143 gmlillicisidryyvaivqavsahrhravllisklscvgilvatvisipellysdlqrs 202
QY 181 SG--IAICTWVYPSDESTKLKSAVLTKV---ILGFFLPFVVMACCVYTIHTLIQAKS 235
DB 203 sseqamrcsli-----tehveafitigvaqmvigfvlplamsfcylviirtllqarf 256
QY 236 SKHKALKVTITVLTVFVLSOPPYNCILLVOTIDAYAMFISNCAVSTNIDICFOVTOTIAF 295
DB 257 ernkaikviiavvvfivqlpyngvvlactvanfnitsstscelskqlnlaydvtyslac 316
QY 296 FHSCNLPVLYVFGFRPRDLVTKLNLGICISQ---AQWVSFTRREGSLKSSMLLE 349
DB 317 vrccvnpflyafgkvkfrndlkfklgclsgqlqswscrh----irssmsve 369

AT 12

AA80114
ID AAG80114 standard; protein; 378 AA.

XX
AC AAG80114;

XX 17-JAN-2002 (first entry)

DE Human CCR7 protein.

KW Chemokine; tumour diagnosis; colorectal; prostatic; organ rejection;
KW inflammation; autoimmune disease; metastasis; bronchial asthma; lupus;
KW chronic bowel inflammation; rheumatoid arthritis; cytostatic;
KW antiinflammatory; antiasthmatic; immunosuppressive; dermatological;
KW antirheumatic; antiarthritic.

OS Homo sapiens.

XX
PN W0200172830-A2.

XX 04-OCT-2001.

XX 02-APR-2001; 2001WO-EP03708.

XX

PR 31-MAR-2000; 2000DE-1016013.

XX (IPFP-) IPF PHARM GMBH.
PA (TORS/) FORSSMANN U.

XX Forssmann W, Adermann K, Heitland A, Spodsberg N;

PI WPI; 2001-626256/72.

XX Diagnostic agent containing two or more receptor-specific ligands,
PT useful for detecting tumors, inflammation etc., also therapeutic use of
PT ligand inhibitors

XX Disclosure; Page 10; 26pp; German.

XX This invention describes a novel diagnostic agent (A) comprising at least
CC two different ligands (I) for receptors (II) that are implicated in
CC disease. (A) are used for the diagnosis of tumors (especially colorectal
CC or prostatic), organ rejection, inflammation and autoimmune diseases.
CC Also inhibitors of (I) are used therapeutically against tumors (and their
CC metastases), inflammation (particularly bronchial asthma or chronic bowel
CC inflammation), or autoimmune diseases (rheumatoid arthritis or lupus),
CC where the (cardio)vascular, lymphatic, respiratory, nervous, digestive,
CC endocrine, motor or urogenital systems or skin are affected, and bone
CC marrow diseases. The products of the invention are chemokine derivatives
CC which have cytostatic, antiinflammatory, antirheumatic, antiarthritic,
CC immunosuppressive, dermatological, antitumor, antiasthmatic,
CC Chemokines act on specific tumor and inflammatory cells through a
CC constellation of chemokine receptors (CR) which control migration and
CC proliferation of these cells. AAG80045-AAG80128 represent human chemokine
CC fragments used to illustrate the method of the invention.

XX Sequence 378 AA;

Query Match 42.1%; Score 780.5; DB 22; Length 378;
Best Local Similarity 42.9%; Pred. No. 1.2e-75;
Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

QY 1 MADDYGESESSMEDYVNFDFCYCKNNVRQFASHFLPPLYWLVFIVGALGNSLVILV 60
DB 28 vtdydgntt-----vdytlfeslcskdkvrnfkawlplmysislfcvlgilnglvlt 82
QY 61 YWYTRVKTMTDFLLNLAIADLLFLVLPFWATAAADQMKFQFMCKVNSYKMFYS 120
DB 83 yiyfkrktmtdtyllnlavadiilflitlpfwaysaakswvfgvhfcklifaikmsffs 142
QY 121 CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKMVCFTIWLAAALCIPILYSQIKEE 180
DB 143 gmlillicisidryyvaivqavsahrhravllisklscvgilvatvisipellysdlqrs 202
QY 181 SG--IAICTWVYPSDESTKLKSAVLTKV---ILGFFLPFVVMACCVYTIHTLIQAKS 235
DB 203 sseqamrcsli-----tehveafitigvaqmvigfvlplamsfcylviirtllqarf 256
QY 236 SKHKALKVTITVLTVFVLSOPPYNCILLVOTIDAYAMFISNCAVSTNIDICFOVTOTIAF 295
DB 257 ernkaikviiavvvfivqlpyngvvlactvanfnitsstscelskqlnlaydvtyslac 316
QY 296 FHSCNLPVLYVFGFRPRDLVTKLNLGICISQ---AQWVSFTRREGSLKSSMLLE 349
DB 317 vrccvnpflyafgkvkfrndlkfklgclsgqlqswscrh----irssmsve 369

RESULT 13

AA80114

ID AAB50859 standard; protein; 378 AA.

XX
AC AAB50859;

XX 16-MAR-2001 (first entry)

XX Human CCR7.

XX

XX Human; chemokine receptor 7; CCR7; chemokine beta-9; Ckbeta-9;
 KW allergy; autoimmune disease; ischaemia; atherosclerosis; cancer;
 KW chronic inflammatory disorder; organ transplant; tissue graft;
 KW chronic myelogenous leukaemia; infection.
 XX Homo sapiens.
 XX OS
 XX PN US6153441-A.
 XX PD 28-NOV-2000.
 XX XX
 XX PF 17-FEB-1999; 99US-0251545.
 XX XX
 XX PR 17-FEB-1998; 98US-0074883.
 XX XX
 XX PA (SMIK) SMITHKLINE BEECHAM CORP.
 XX XX
 XX PI Appelbaum ER, White JR, Sarau HM;
 XX XX
 XX XX I; 2001-049151/06.
 XX XX
 XX PT Identifying agonists or antagonists of interaction between human
 PT protein, chemokine beta-9 and human CC chemokine receptor 7, by
 PT contacting cell expressing receptor with test compound -
 XX XX
 XX PS Claim 1; Fig 1; 20pp; English.
 XX XX
 XX CC The present sequence is human chemokine receptor 7 (CCR7), a cellular
 CC receptor for chemokine beta-9 (Ckbeta-9). The sequence may be
 CC used in a method for discovering agonists and antagonists of the
 CC interaction between Ckbeta-9 and CCR7. A cell expressing CCR7
 CC polypeptide on its surface, associated with a component capable of
 CC providing a detectable signal in response to binding of Ckbeta-9, is
 CC contacted with a compound in the presence of labelled or unlabelled
 CC Ckbeta-9. The compound is identified as an agonist/antagonist by
 CC determining whether it activates or inhibits the detectable signal.
 CC The method is useful for identifying agonists and antagonists of the
 CC interaction between Ckbeta-9 and CCR7 which are useful for treating
 CC diseases including allergic disorders, autoimmune diseases,
 CC ischaemia/perfusion injury, development of atherosclerotic plaques,
 CC cancer, chronic inflammatory disorders, chronic rejection of
 CC transplanted organs or tissue grafts, chronic myelogenous leukaemia, and
 CC infection by HIV and other pathogens.
 XX XX
 XX SQ Sequence 378 AA;

Query Match 42.1%; Score 780.5; DB 22; Length 378;
 Best Local Similarity 42.9%; Pred. No. 1.2e-75;
 Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

QY 1 MADDYGESTSSMEDYVNFNFTDFYCEKNVROFASHFLPPLVWLVFIVGALGNSLVLY 60
 Db 28 vtdyigdnnt-----vdytlfeslcskdkvrfkawlplmlysiicfcvgllnglvlt 82
 QY 61 YWCTRVKVTMTDFLLNLATADLLFLVLPFWATAADQWKQFTFMCKVNSMYKMFYS 120
 Db 83 yifkrlktmtdtyllnlavadihlitlpfwaysaakswvgvfhcklifaikmsffs 142
 QY 121 CVLLIMCISVDRIYIAQAQRAHTRKRLLYSKMVCFTIWLAAALCIPILYSQKEE 180
 Db 143 gmlilicisidryvaivqavsahrhrarvilliskscvgiwtlatvisipellysdigr 202
 QY 181 SG--IAICTWVPSPDESKLKSAVLTUKV---ILGFPLFPVWVWACCYTIHILIOAKKS 235
 Db 203 sseqamrcsl-----tehveafitigvaqmvigfivpllamscfylviirtlqarf 256
 QY 236 SKHAKLVTTVLTVFVLSQPPYNCILLVQTIDAYAMFISNCVASTNIDICFQVOTIAF 295
 Db 257 ernkaikviiavvvfivqipngvvaqgtvanfnststcelskqlniaydvtyslac 316
 QY 296 FHSLNPNVLYVFGERRFRDLVTKNLKNGCISQ---AQWVSFTRREGSLKLSMLLE 349

Db 317 vrcvnpfyafyigvfrndflkfldgcisqelqrwsscrh-----irssmsve 369
 RESULT 14
 AAR53743
 ID AAR53743 standard; Protein; 410 AA.
 XX AC AAR53743;
 XX DT 02-FEB-1995 (first entry)
 XX DE Putative seven transmembrane receptor (V31).
 XX KW Primer; seven transmembrane receptor; receptor; amplification; PCR;
 KW polymerase chain reaction.
 XX OS Homo sapiens.
 XX PN WO9412635-A.
 XX PD 09-JUN-1994.
 XX PF 17-NOV-1993; 93WO-US111153.
 XX PR 17-NOV-1992; 92US-0977452.
 XX PA (ICOS-) ICOS CORP.
 XX PI Godiska R, Gray PW, Schweickart VL;
 XX WPI; 1994-200264/24.
 DR N-PSDB; AAO66153.
 XX DNA encoding seven transmembrane receptors - used to develop
 PT prods, for use as therapeutic or diagnostic agents for conditions
 PT involving the receptors.
 XX Example 2; Page 46-48; 100pp; English.
 XX Two primers (AAO66148, AAO66149) were used to amplify human genomic DNA
 CC purified from leukocytes. Approximately 1000 clones were isolated
 CC after the initial amplification reaction and probed with sequences
 CC specific for seven transmembrane receptors IL8R1, ATR2R and R20.
 CC Clones which did not hybridise were then chosen for sequence
 CC analysis. Three new clones were identified that appeared to encode
 CC seven transmembrane receptor segments. Two more primers (AAO66151,
 CC AAO66152) were used to isolate a full length version of one of these
 CC clones, one of which was designated V31 and encoded this
 CC polypeptide.
 XX SQ Sequence 410 AA;

Query Match 42.1%; Score 780.5; DB 15; Length 410;
 Best Local Similarity 42.9%; Pred. No. 1.4e-75;
 Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

QY 1 MADDYGESTSSMEDYVNFNFTDFYCEKNVROFASHFLPPLVWLVFIVGALGNSLVLY 60
 Db 60 vtdyigdnnt-----vdytlfeslcskdkvrfkawlplmlysiicfcvgllnglvlt 114
 QY 61 YWCTRVKVTMTDFLLNLATADLLFLVLPFWATAADQWKQFTFMCKVNSMYKMFYS 120
 Db 115 yifkrlktmtdtyllnlavadihlitlpfwaysaakswvgvfhcklifaikmsffs 174
 QY 121 CVLLIMCISVDRIYIAQAQRAHTRKRLLYSKMVCFTIWLAAALCIPILYSQKEE 180
 Db 175 gmlilicisidryvaivqavsahrhrarvilliskscvgiwtlatvisipellysdigr 234
 QY 181 SG--IAICTWVPSPDESKLKSAVLTUKV---ILGFPLFPVWVWACCYTIHILIOAKKS 235
 Db 235 sseqamrcsl-----tehveafitigvaqmvigfivpllamscfylviirtlqarf 288

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OM protein - protein search, using sw model

Run on: October 1, 2002, 06:26:10 ; Search time 12.97 Seconds
(without alignments)
672.316 Million cell updates/sec

Title: US-09-522-752-2
Perfect score: 1854
Sequence: 1 MADDYGESTSSMEDYVNFN.....EGSLKSSMLLETTSGALS 357

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 231628 seqs, 24425594 residues

Total number of hits satisfying chosen parameters: 231628

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents.AA:*
1: /cgn2_6/ptodata/2/1aa/5A.COMB.pep:*
2: /cgn2_6/ptodata/2/1aa/5B.COMB.pep:*
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4: /cgn2_6/ptodata/2/1aa/6B.COMB.pep:*
5: /cgn2_6/ptodata/2/1aa/PCTUS.COMB.pep:*
6: /cgn2_6/ptodata/2/1aa/backfiles1.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1854	100.0	357	4	US-09-266-464-2
2	780.5	42.1	358	1	US-08-153-848-19
3	780.5	42.1	358	3	US-09-299-843A-19
4	780.5	42.1	358	4	US-09-088-337B-19
5	780.5	42.1	358	5	PCT-US93-11153-19
6	780.5	42.1	378	1	US-08-153-848-15
7	780.5	42.1	378	3	US-09-299-843A-15
8	780.5	42.1	378	4	US-09-251-545-1
9	780.5	42.1	378	4	US-09-088-337B-15
10	780.5	42.1	378	5	PCT-US93-11153-15
11	780.5	42.1	410	1	US-08-153-848-7
12	780.5	42.1	410	3	US-09-299-843A-7
13	780.5	42.1	410	4	US-09-088-337B-7
14	780.5	42.1	410	5	PCT-US93-11153-7
15	758.5	40.9	378	1	US-08-383-751A-2
16	758.5	40.9	378	3	US-08-352-678-2
17	758.5	40.9	378	4	US-09-045-583-49
18	758.5	40.9	378	5	PCT-US93-09636-2
19	757.5	40.9	378	3	US-09-299-843A-66
20	757.5	40.9	378	4	US-09-088-337B-66
21	757.5	40.9	378	4	US-08-153-848-24
22	742	40.0	359	1	US-08-153-848-24
23	742	40.0	359	3	US-09-299-843A-24
24	742	40.0	359	4	US-09-088-337B-24
25	742	40.0	359	5	PCT-US93-11153-24
26	721	38.9	361	2	US-08-902-294-2
27	721	38.9	361	3	US-09-178-637-2

28	652	35.2	350	2	US-08-966-316-18	Sequence 18, Appl
29	638.5	34.4	374	4	US-09-045-583-48	Sequence 48, Appl
30	637	34.4	350	2	US-08-966-316-16	Sequence 16, Appl
31	624	33.7	342	4	US-09-116-498-6	Sequence 6, Appl
32	619	33.4	342	4	US-09-116-498-4	Sequence 4, Appl
33	612	33.0	342	2	US-08-742-011-2	Sequence 2, Appl
34	612	33.0	342	4	US-09-275-384B-5	Sequence 5, Appl
35	612	33.0	342	4	US-09-116-498-2	Sequence 2, Appl
36	612	33.0	342	4	US-09-449-437A-2	Sequence 2, Appl
37	598	32.3	352	4	US-09-045-583-52	Sequence 52, Appl
38	596	32.1	352	4	US-09-087-232A-13	Sequence 13, Appl
39	596	32.1	352	4	US-08-861-105-14	Sequence 14, Appl
40	596	32.1	352	4	US-08-575-967A-2	Sequence 2, Appl
41	590	31.8	352	3	US-08-466-343D-2	Sequence 2, Appl
42	585.5	31.6	360	4	US-08-875-573-20	Sequence 20, Appl
43	585.5	31.6	360	4	US-09-232-878-2	Sequence 2, Appl
44	585.5	31.6	360	4	US-09-045-583-55	Sequence 55, Appl
45	582	31.4	360	1	US-08-202-056-7	Sequence 7, Appl

ALIGNMENTS

RESULT 1
US-09-266-464-2
; Sequence 2, Application US/09266464
; GENERAL INFORMATION:
; APPLICANT: Andrew, David P.
; APPLICANT: Zabel, Brian A.
; APPLICANT: Ponath, Paul D.
; TITLE OF INVENTION: ANTI-GPR-9-6 ANTIBODIES AND METHODS OF IDENTIFYING AGENTS WHICH MODULATE GPR-9-6 FUNCTION
; FILE REFERENCE: LKS98-16
; CURRENT APPLICATION NUMBER: US/09/266.464
; CURRENT FILING DATE: 1999-03-11
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 357
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-266-464-2

Query Match	100.0%	Score	1854;	DB	4;	Length	357;
Best Local Similarity	100.0%	Pred. No.	7.8e-162;				
Matches	357;	Conservative	0;	Mismatches	0;	Indels	0;
Qy	1	MADDYGESTSSMEDYVNFNFTDFYCEKNNVQFASHFLPPLYWLVFIVGALGNSLVILV	60				
Db	1	MADDYGESTSSMEDYVNFNFTDFYCEKNNVQFASHFLPPLYWLVFIVGALGNSLVILV	60				
Qy	61	YWCTRVKTMDFLNLAIADLLFLVLPFWAIAAADOWKQFOTFCMKVNSMYKMFYS	120				
Db	61	YWCTRVKTMDFLNLAIADLLFLVLPFWAIAAADOWKQFOTFCMKVNSMYKMFYS	120				
Qy	121	CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKWVCFTIWLAAALCIPEILYSQIKEE	180				
Db	121	CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKWVCFTIWLAAALCIPEILYSQIKEE	180				
Qy	181	SGIAICTWYPSDESTKLKSAVLTILKVLGFFLPFWMACCVYTIHHTLIQAKSKHKA	240				
Db	181	SGIAICTWYPSDESTKLKSAVLTILKVLGFFLPFWMACCVYTIHHTLIQAKSKHKA	240				
Qy	241	LKVTITVLTVFVLSOFFPNCILLVQTIDAYAMFISNCAVSTNIDICFQVQTQIAFFHSL	300				
Db	241	LKVTITVLTVFVLSOFFPNCILLVQTIDAYAMFISNCAVSTNIDICFQVQTQIAFFHSL	300				
Qy	301	NPVLVYFVGERRDLVKTLKNLGICISQAOVSWFTRREGSLKSSMLLETTSGALS	357				
Db	301	NPVLVYFVGERRDLVKTLKNLGICISQAOVSWFTRREGSLKSSMLLETTSGALS	357				

[illegible]

```

Db      297 VRCVNPFLYAFYGVFRNDLFLKFLDXDGLCLSQLEQLQWSSCRH-----IIRSSMSVE 349

RESULT          3
US-09-299-843A-19
; Sequence 19, Application US/09299843A
; Patent No. 6107475
; GENERAL INFORMATION:
; APPLICANT: Godiska, Ronald
; APPLICANT: Gray, Patrick W.
; APPLICANT: Schweikart, Vicki L.
; TITLE OF INVENTION: No. 6107475el Seven Transmembrane Receptors
; NUMBER OF SEQUENCES: 66
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
; ADDRESSEE: Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatenIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/299,843A
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 09/088,337
; FILING DATE: 01-JUN-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/153,848
; FILING DATE: 17-NOV-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/977,452
; FILING DATE: 17-NOV-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Jill E. Uhl
; REGISTRATION NUMBER: 43,213
; REFERENCE/DOCKET NUMBER: 27866/32059B
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (312) 474-6300
; TELEFAX: (312) 474-0448
; TELEX:
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 358 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-09-299-843A-19

Query Match           42.1%; Score 780.5; DB 3; Length 358;
Best Local Similarity 42.9%; Pred. No. 8.8e-64;
Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps

Qy      1 MADDYGSEPTSMEDYVNFTDFCYCKNNVRQFAFHLPPLYLVLVFIVGALGNLSILVLV 60
       :|||::||:||||:::||:||||:||||:||||:||||:||||:||||:||||:||||:
Db      8 VTDDYTGDNTT-----VDYTLPSLCSKDKDVRFNKAFLPIMFIYCICFVGLLGNGLVLT 62
       :|||::||:||||:::||:||||:||||:||||:||||:||||:||||:||||:||||:

Qy      61 YWCYTRVKTMDFMLNLAIATDLFLTVPFWIAAAADQWKQTFCMKVVNSMYKNMFSYS 120
       |||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||:
Db      63 YIVFKRLKTMTDTYLLNLAVADLFLLTPFEWAYSAKSWMFVGCHFKLIFFAYIKMFSPS. 122
       :|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||:

Qy      121 CVLLIMCISVDRIATAQAAMRAHTREKRLLLYSKVCFFITWIWLAAALCIPEILYSOIKEE 180
       :|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||:
Db      123 GMILLJCISIDRVVAIQVASIIHRRHARVLLISKLSCVGIWILTAVLSIPELLYSOLQRS 182
       :|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||:

Qy      181 SG--IAICTMVYPDSDESTLKLSAVLTALKV---ILGFPLFPVMACCTYTIITHLIOAKKS 233

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Db      297   VRCVNPFLYAFTGVFRNDLFLKLFKDLCGLSQLEQLQWSSCRH-----IIRSSMSVE    349


RESULT          3
US-09-299-843A-19
; Sequence 19, Application US/09299843A
; Patent No. 6107475
; GENERAL INFORMATION:
; APPLICANT: Godiska, Ronald
; APPLICANT: Gray, Patrick W.
; APPLICANT: Schweikart, Vicki L.
; TITLE OF INVENTION: NO. 6107475el Seven Transmembrane Receptors
; NUMBER OF SEQUENCES: 66
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
; ADDRESSEE: Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/299,843A
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 09/088,337
; FILING DATE: 01-JUN-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/153,848
; FILING DATE: 17-NOV-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/977,452
; FILING DATE: 17-NOV-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Jill E. Uhl
; REGISTRATION NUMBER: 43,213
; REFERENCE/DOCKET NUMBER: 27866/32059B
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (312) 474-6300
; TELEFAX: (312) 474-0448
; TELEX:
; INFORMATION FOR SEQ ID NO: 19:
;     SEQUENCE CHARACTERISTICS:
;         LENGTH: 358 amino acids
;         TYPE: amino acid
;         TOPOLOGY: linear
;     MOLECULE TYPE: protein
US-09-299-843A-19



Query Match       42.1%; Score 780.5; DB 3; Length 358;
Best Local Similarity 42.9%; Pred. No. 8.e-64;
Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps

Qy      1  MADDYGESESSMEDYYNFFTDIFYCEKNNVROFAHFPLPPLYLVFIYGALGNSILVLV    60
        :|||:::||:||||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||
Db      8  VTDDYTGDNTT-----VDYTLPSLSCLKDKDVRNFKAFLPIMIFSIICTFVGLLGNGLVLT    62
        ||| |
Qy      61  YWCYTRVKMTDMFLNLAIATDLFLTVPFWIAAAADQWKQTFCMKVVNSMYKMNFYS    120
        ||| |
Db      63  YIVFKRLKTMDTYLLNLAVADILELLTPFEWAYSAAKSWVFCHVCKLIIFAITYKMSPFS.    122
        ||| |
Qy      121  CVLLIMCISVDRIATAQAAMRAHTREKRLLLTSKWVCFTIWLAALCALCIPEILYSOIKEE    180
        :|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||
Db      123  GMILLJCISIDRVAIQVASIIHRHHRRARVLLISKSCVGWIWLATVLSIPELLYSOLQRS    182
        ||| |
Qy      181  SG--IAICTMVYPDESSTKLKSAVLTKV---ILGFPLFPVMACCYTTIIHTLIOAKKS    235
```

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 1, 2002, 06:26:30 ; Search time 46.91 Seconds
(without alignments)
731.270 Million cell updates/sec

Title: US-09-522-752-2
Perfect score: 1854
Sequence: 1 MADDYGSESTSMEDYVNFN.....EGSLKLSMLETTSGLSL 357

Scoring table:
BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283138 seqs, 96089334 residues

To: Number of hits satisfying chosen parameters: 283138

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR_71.*
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	780.5	42.1	378	2 B55735	lymphocyte-specific
2	758.5	40.9	378	2 A45680	G protein-coupled
3	757.5	40.9	378	2 A55735	G protein-coupled
4	652	35.2	350	2 JN0621	G protein-coupled
5	638.5	34.4	369	2 JC5068	G protein-coupled
6	596	32.1	352	2 A43113	chemokine (C-C) re
7	585.5	31.6	360	2 A57160	chemokine (C-C) re
8	582	31.4	360	2 A53611	interleukin-8 rece
9	577	31.1	360	2 JC4587	chemokine (C-C) re
10	575	31.0	360	2 JC2443	chemokine (C-C) re
11	571	30.8	355	2 A45177	chemokine (C-C) re
12	569	30.7	374	2 I38450	chemokine (C-C) re
13	569	30.7	383	2 S55594	G protein-coupled
14	566	30.5	358	2 A53752	interleukin-8 rece
15	565.5	30.5	353	2 S28787	neuropeptide Y/pep
16	560	30.2	355	2 G02436	chemokine (C-C) re
17	558.5	30.1	350	2 A39445	interleukin-8 rece
18	558.5	30.1	355	2 J01231	interleukin-8 rece
19	557.5	30.1	352	2 A45747	neuropeptide Y/pep
20	555	29.9	359	2 I49341	MIP-1 alpha recept
21	547.5	29.5	352	2 G00048	fusin (LESTRA) - c
22	545	29.4	359	2 A48921	interleukin-8 rece
23	543.5	29.3	355	2 S42096	interleukin-8 rece
24	528	28.5	374	2 S32785	G protein-coupled
25	527.5	28.5	355	2 I49339	macrophage inflam
26	526.5	28.4	374	2 S42628	G protein-coupled
27	524	28.3	327	2 S56162	MBP15 protein - h
28	521.5	28.1	359	2 S15403	angiotensin II rec
29	521.5	28.1	372	2 S26667	G protein-coupled

30	517.5	27.9	359	2 S44425	angiotensin II rec
31	516.5	27.9	359	2 JC1104	angiotensin II rec
32	512	27.6	355	2 JC5067	G protein-coupled
33	512	27.6	367	2 JE0349	interferon-inducib
34	508.5	27.4	359	2 A48857	angiotensin II rec
35	505.5	27.3	359	2 A42656	angiotensin II rec
36	504.5	27.2	359	2 I39418	angiotensin II rec
37	504.5	27.2	359	2 JC2134	angiotensin II rec
38	501.5	27.0	359	2 JC1194	angiotensin II rec
39	500	27.0	356	2 I49340	MIP-1 alpha recept
40	498.5	26.9	359	2 JH0621	angiotensin II rec
41	494.5	26.7	359	2 JQ1516	angiotensin II rec
42	491.5	26.5	362	2 JN0694	interleukin-8 rece
43	483	26.1	354	2 A23669	orphan G protein-c
44	482.5	26.0	355	2 JC4304	G protein-coupled
45	479	25.8	354	2 B55733	

ALIGNMENTS

RESULT 1

B55735
lymphocyte-specific G protein-coupled receptor EB11 - human
N;Alternate names: Burkitt's lymphoma receptor 2; Epstein-Barr virus induced protein
C;Species: Homo sapiens (man)
C;Date: 07-Jul-1995 #sequence_revision 07-Jul-1995 #text_change 19-May-2000
C;Accession: B55735; S52443
R;Schweickart, V.L.; Raport, C.J.; Godiska, R.; Byers, M.G.; Eddy Jr., R.L.; Shows, T
Genomics 23, 643-650, 1994
A;Title: Cloning of human and mouse EB11, a lymphoid-specific G-protein-coupled recep
A;Reference number: A55735; MUID:95154835
A;Accession: B55735
A;Status: preliminary
A;Molecule type: mRNA
A;Residues: 1-378 <SCH>
A;Cross-references: GB:L31581; NID:g468319; PIDN:AAA74231.1; PID:g468320
R;Burgstahler, R.; Kempkes, B.; Staube, K.; Lipp, M.
submitted to the EMBL Data Library, February 1995
A;Description: The expression of the chemokine receptor BLR2/EB11 is specifically tra
A;Reference number: S52443
A;Accession: S52443
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 21-378 <BUR>
A;Cross-references: EMBL:X84702
C;Genetics:
A;Gene: GDB:CMKBR7; EB11; BLR2; CCR7
A;Cross-references: GDB:342065; OMIM:600242
A;Map position: 17q12-17q21.2
C;Superfamily: vertebrate rhodopsin
C;Keywords: G protein-coupled receptor

Query Match 42.1%; Score 780.5; DB 2; Length 378;
Best Local Similarity 42.9%; Pred. No. 3.9e-60;
Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

QY	1	MADDYGSESTSMEDYVNFNFDYCEKNVVRQFASHFLPPLVWLVFVAGALNSILVY	60
DB	28	VTDDYIGDNTT-----VDYTLFSLCSKCKDVRNFKAWFLPIMYSIICIFVGLLGNLVLT	82
QY	61	YWYCTRVKTMDFLLNLATADLLFLVLPFWATAADONKQFQFMCKVNSMYKMFYS	120
DB	83	YIYFKRLKTTDTYLLNLAVADILFLTLTLPFWAYSAAKSWVGFVHFKLFAIKMFSS	142
QY	121	CVLLIMCISVDVRYIAQAAMRAHREKRLLYSKMVCFTIWLAAALCIPELYSQKEE	180
DB	143	GMLLLCISIDRYVAIVQVSAHRARVLLISLSCVGIWILATVLSIPELLYSDLQRS	202
QY	181	SG--IAICTWYPSDESTKLKSAVLTLLK-----ILGFFLPFVVMACCYTIITHTLIQAKS	235
DB	203	SSEQAMRCSLI-----TEHVEAFITIQVQMVIGFLVPLPLAMSFCLVIIRTLQARNF	256

A: Residues: 1-378 <SCH>
 A: Cross-references: GB:I31580; NID:g468340; PIDN:AAA74232.1; PID:g468341
 C: Superfamily: vertebrate rhodopsin
 C: Keywords: G protein-coupled receptor

Query Match 40.9%; Score 757.5; DB 2; Length 378;
Best Local Similarity 43.2%; Pred. No. 3.8e-58;
Matches 156; Conservative 74; Mismatches 108; Indels 23; Gaps

[illegible]

203 SGEDFLRCSLSAQVE-----ALITIQVAQWVFGFLVPMAMSFCYLLIRTLQARNF 250

QY 236 FHSCLNPLVIVFGKRTKRDGKVLKLNAGGCLSSQ ---WSEFLR-KDGSUNKSSMDELLI 370

Db 317 VRCCVNPFLYAFIGVKERSDLFLKFLKDGLGCLSQERLHWSSCRHVRNASV---SMEAEIT 373

QV 352 S 352

RESULT 4
JN0621
G protein-coupled receptor type B - bovine
C:Species: Bos primigenius taurus (cattle)

C:Species: *Sos plimigenus* caurus (Cactili)
C:Date: 24-Feb-1994 #sequence_revision 24-Feb-1994 #text_change 19-May-2000
R:Accession: JN0621
R:Matsuoka, I.; Mori, T.; Aoki, J.; Sato, T.; Kurihara, K.
Biochem. Biophys. Res. Commun. 194, 504-511, 1993
A:Title: Identification of novel members of G-protein coupled receptor superfamily ex
A:Reference number: JN0621; MUID:93326166
A:Accession: JN0621

A: Molecule type: *hikrka*
A: Residues: 1-350 (MAT)
A: Cross-references: GB:563848; NID:g399710; PIDN:AAB27547.1; PID:g399711
A: Experimental source: Tongue taste papillae
C: Comment: This protein is involved in modulating taste sensitivity or regeneration of taste buds
C: Superfamily: vertebrate rhodopsin
C: Keywords: G protein-coupled receptor; glycoprotein; receptor; transmembrane protein
F: 42-66/Domain: transmembrane #status predicted <TM1>
F: 80-99/Domain: transmembrane #status predicted <TM2>
F: 114-135/Domain: transmembrane #status predicted <TM3>
F: 154-175/Domain: transmembrane #status predicted <TM4>
F: 200-222/Domain: transmembrane #status predicted <TM5>
F: 242-265/Domain: transmembrane #status predicted <TM6>
F: 284-306/Domain: transmembrane #status predicted <TM7>
F: 6.19/Binding site: carboxylate (Asn) (covalent) #status predicted

```
Query Match      35.2%; Score 652; DB 2; Length 350;
Best Local Similarity 37.9%; Pred. No. 5e-49;
Matches 127; Conservative 70; Mismatches 114; Indels 24; Gaps
1 MADDYGSEST-----SSMEDYVNFNTDFYCEKNVRQFASHEFLPYLVLFIVGALGN 54
```


GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 1, 2002, 06:29:56 ; Search time 13.53 Seconds
(without alignments)
1021.647 Million cell updates/sec

Title: US-09-522-752-2

Perfect score: 1854

Sequence: 1 MADDYGSSTSMEDYVNFN.....EGSLKLSMLTTGALSLSL 357

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 105224 seqs, 38719550 residues

To: Number of hits satisfying chosen parameters: 105224

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_40.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Match	Length	DB ID	Description
1	1854	100.0	357	1 CKR9_HUMAN
2	1642	88.6	369	1 CKR9_MOUSE
3	780.5	42.1	378	1 CKR7_HUMAN
4	757.5	40.9	378	1 CKR7_MOUSE
5	652	35.2	350	1 CKRB_BOVIN
6	638.5	34.4	374	1 CKRB_MOUSE
7	637	34.4	350	1 CKRB_HUMAN
8	624	33.7	342	1 CCR6_MACNE
9	619	33.4	342	1 CCR6_CERAE
10	617	33.3	343	1 CCR6_MACMU
11	612	33.0	342	1 CCR6_HUMAN
12	606	32.7	352	1 CKR5_CERTO
13	605	32.6	367	1 CKR6_MOUSE
14	604	32.6	352	1 CKR5_CERAE
15	604	32.6	352	1 CKR5_GORGO
16	603	32.5	352	1 CKR5_MACMU
17	602	32.5	352	1 CKR5_PAPHA
18	601	32.4	352	1 CKR5_PONPY
19	598	32.3	352	1 CKR5_PANTR
20	598	32.3	352	1 CKR5_TRAFR
21	598	32.3	352	1 CKR5_TRAPH
22	597	32.2	352	1 CKR5_PYGBI
23	597	32.2	352	1 CKR5_PYGNE
24	596	32.1	352	1 CKR5_HUMAN
25	595	32.1	352	1 CKR5_HYLLE
26	585.5	31.6	354	1 CKR5_MOUSE
27	585.5	31.6	360	1 CKR4_HUMAN
28	582	31.4	360	1 IL8B_HUMAN
29	578	31.2	373	1 CKR2_RAT
30	577	31.1	360	1 CKR4_MOUSE
31	575	31.0	373	1 CKR2_MOUSE
32	573.5	30.9	354	1 CKR5_RAT
33	573	30.9	360	1 CKR2_MACMU

ALIGNMENTS

RESULT 1.

ID	CKR9_HUMAN	STANDARD	PRT	357 AA.
AC	P51686;			
DT	01-OCT-1996 (Rel. 34, Created)			
DT	01-OCT-1996 (Rel. 34, Last sequence update)			
DT	01-MAR-2002 (Rel. 41, Last annotation update)			
DE	C-C chemokine receptor type 9 (C-C CKR-9) (CCR-9) (GPR-9-6).			
DE	6).			
GN	CCR9 OR CMKBR9.			
OS	Homo sapiens (Human).			
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;			
OC	Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.			
OX	NCBI_TaxID=9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RA	Lautens L.L., Tiffany H.L., Gao J.-L., Modi W., Murphy P.M.,			
RA	Bonner T.I.;			
RL	Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.			
RN	[2]			
RP	CHARACTERIZATION.			
RX	MEDLINE=99248133; PubMed=10229797;			
RA	Zaballos A., Gutierrez J., Varona R., Ardavin C., Marquez G.;			
RT	"Cutting edge: Identification of the orphan chemokine receptor GPR-9-6 as CCR9, the receptor for the chemokine TECK.";			
RL	J. Immunol. 162:5671-5675(1999).			
CC	-I- FUNCTION: Receptor for chemokine SCYA25/TECK. Subsequently transduces a signal by increasing the intracellular calcium ions level. Alternative coreceptor with CD4 for HIV-1 infection.			
CC	-I- SUBCELLULAR LOCATION: Integral membrane protein.			
CC	-I- TISSUE SPECIFICITY: HIGHLY EXPRESSED IN THE THYMUS AND LOW IN LYMPH NODES AND SPLEEN.			
CC	-I- SIMILARITY: BELONGS TO FAMILY 1 OF G-PROTEIN COUPLED RECEPTORS.			
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (see http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).			
CC	EMBL: U45982; AAA93319.1; -			
DR	CCRD6; GCR1943; -			
DR	MM; 604738; -			
DR	InterPro: IPR004069; Chemokine9_receptor.			
DR	InterPro: IPR000276; GPCR_Rhodpsn.			
DR	Pfam: PF00001; 7tm.1; 1.			
DR	PRINTS: PR01531; CHEMOKINER9.			
DR	PRINTS: PR02037; GPCRHOOPS.			
DR	PROSITE: PS00237; G_PROTEIN_RECEP_F1_1; 1.			
DR	PROSITE: PS00262; G_PROTEIN_RECEP_F1_2; 1.			
KW	G-protein coupled receptor; Transmembrane; Glycoprotein.			
FT	DOMAIN 1 37 EXTRACELLULAR (POTENTIAL).			
FT	TRANSMEM 38 64 1 (POTENTIAL).			
FT	DOMAIN 65 73 CYTOPLASMIC (POTENTIAL).			

P32246 homo sapien
Q28807 pan troglod
P56498 felis silve
P41597 homo sapien
Q28519 macaca mula
P35344 oryctolagus
P25930 bos taurus
Q28422 gorilla gor
P55919 gorilla gor
P51677 homo sapien
P55920 pan troglod
P21109 oryctolagus

34 571 30.8 355 1 CKR1_HUMAN
35 570.5 30.8 353 1 IL8B_PANTR
36 569.5 30.7 353 1 CCR4_FELCA
37 569 30.7 374 1 CKR2_HUMAN
38 568.5 30.7 353 1 IL8B_MACMU
39 566 30.5 358 1 IL8B_BOVIN
40 565.5 30.5 353 1 CCR4_BOVIN
41 565.5 30.5 353 1 IL8B_GORGO
42 564.5 30.4 350 1 IL8A_GORGO
43 563 30.4 355 1 CKR3_HUMAN
44 558.5 30.1 350 1 IL8A_PANTR
45 558.5 30.1 355 1 IL8A_RABIT

```

FT TRANSMEM 74 94 2 (POTENTIAL).
FT DOMAIN 95 108 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 109 130 3 (POTENTIAL).
FT DOMAIN 131 148 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 149 169 4 (POTENTIAL).
FT DOMAIN 170 198 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 199 226 5 (POTENTIAL).
FT DOMAIN 227 242 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 243 268 6 (POTENTIAL).
FT DOMAIN 269 292 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 293 310 7 (POTENTIAL).
FT DOMAIN 311 357 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 20 186 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT DISULFID 107 186 BY SIMILARITY.
SQ SEQUENCE 357 AA; 40713 MW; 96982E0B922F6B31 CRC64;

Query Match 100.0%; Score 1854; DB 1; Length 357;
Best Local Similarity 100.0%; Pred. No. 6.3e-115; Indels 0; Gaps 0;
Matches 357; Conservative 0; Mismatches 0;

1 MADDYSESTSMEDYVNFNFTDFYCEKNNVRQFASHPPLPLYWLVFVIGALNSLVILV 60
|||||
1 MADDYSESTSMEDYVNFNFTDFYCEKNNVRQFASHPPLPLYWLVFVIGALNSLVILV 60
Db

61 YWYCTRVKVTMTDMFLNLAIADLLFLVLPFWATAAADQWKFQTFMCKVNSMYKMFYS 120
61 YWYCTRVKVTMTDMFLNLAIADLLFLVLPFWATAAADQWKFQTFMCKVNSMYKMFYS 120
Db

121 CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKMVCFTIWLAAALCIPILYSQIKEE 180
121 CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKMVCFTIWLAAALCIPILYSQIKEE 180
Db

181 SGIAICTMVPSPDESTKLKSAVLTKVILGFPLPVVMACCYTIITHTLQAKSSKHKA 240
181 SGIAICTMVPSPDESTKLKSAVLTKVILGFPLPVVMACCYTIITHTLQAKSSKHKA 240
Db

241 LKVTITVTLVFLVSQFPYNCILLVOTIDAYAMFISNCAVSNIDICFQVQTIAFFHSCL 300
241 LKVTITVTLVFLVSQFPYNCILLVOTIDAYAMFISNCAVSNIDICFQVQTIAFFHSCL 300
Db

301 NPVLVYVGERFRRLDLVTKNLGICISQAQWVSFTRREGSKLSSMLLETSGALS 357
301 NPVLVYVGERFRRLDLVTKNLGICISQAQWVSFTRREGSKLSSMLLETSGALS 357
Db

RESULT 2
CKR9_MOUSE STANDARD; PRT; 369 AA.
CKR9_MOUSE
Q9WU7;
30-MAY-2000 (Rel. 39, Created)
30-MAY-2000 (Rel. 39, Last sequence update)
16-OCT-2001 (Rel. 40, Last annotation update)
C-C chemokine receptor type 9 (C-CR-9) (CC-CR-9) (CCR-9)
(Chemokine C-C receptor 10).
CCR9 OR CNKBR10.
Mus musculus (Mouse).
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
NCBI_TaxID=10090;
[1]
SEQUENCE FROM N.A.
TISSUE=Thymus;
MEDLINE=95248139; PubMed=10229797;
Zaballos A., Gutierrez J., Varona R., Ardavin C., Marquez G.;
"Cutting edge: identification of the orphan chemokine receptor GPR-9-6
as CCR9, the receptor for the chemokine TECK";
J. Immunol. 162:5671-5675(1999).
- FUNCTION: RECEPTOR FOR CHEMOKINE SCYA25/TECK. SUBSEQUENTLY
TRANSDUCES A SIGNAL BY INCREASING THE INTRACELLULAR CALCIUM IONS
LEVEL.
- SUBCELLULAR LOCATION: Integral membrane protein.
- TISSUE SPECIFICITY: HIGHLY EXPRESSED IN THE THYMUS AND LOW IN

```

```

CC LYMPH NODES AND SPLEEN.
CC - SIMILARITY: BELONGS TO FAMILY 1 OF G-PROTEIN COUPLED RECEPTORS.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: AJ132336; CAB43480.1; -.
CC MGD: MGI:1341902; Cmkbr10.
CC InterPro: IPR004069; Chemokine9_receptor.
CC InterPro: IPR000276; GPCR_Rhodpsn.
CC Pfam: PF00001; 7tm_1; 1.
CC PRINTS: PR01531; CHEMOKINER9.
CC PRINTS: PR00237; GPCRHHODPSN.
CC PROSITE: PS00237; G_PROTEIN_RECEP_FL_1; 1.
CC PROSITE: PS00262; G_PROTEIN_RECEP_FL_2; 1.
CC G-protein coupled receptor; Transmembrane; Glycoprotein.
CC KW DOMAIN 1 49 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 50 76 1 (POTENTIAL).
FT DOMAIN 77 85 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 86 106 2 (POTENTIAL).
FT DOMAIN 107 120 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 121 142 3 (POTENTIAL).
FT DOMAIN 143 160 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 161 181 4 (POTENTIAL).
FT DOMAIN 182 210 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 211 238 5 (POTENTIAL).
FT DOMAIN 239 254 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 255 280 6 (POTENTIAL).
FT DOMAIN 281 304 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 305 322 7 (POTENTIAL).
FT DOMAIN 323 369 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 32 32 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT DISULFID 119 198 BY SIMILARITY.
SQ SEQUENCE 369 AA; 41913 MW; 6971F7F0A24B4AE CRC64;

Query Match 88.6%; Score 1642; DB 1; Length 369;
Best Local Similarity 86.6%; Pred. No. 4.9e-101;
Matches 309; Conservative 21; Mismatches 27; Indels 0; Gaps 0;

Qy 1 MADDYSESTSMEDYVNFNFTDFYCEKNNVRQFASHPPLPLYWLVFVIGALNSLVILV 60
|||||
13 MFDDFSYDSTASTDDYNNLNFSSFECKNNVRQFASHPPLPLYWLVFVIGALNSLVILV 72
Db

61 YWYCTRVKVTMTDMFLNLAIADLLFLVLPFWATAAADQWKFQTFMCKVNSMYKMFYS 120
|||||
73. YWYCTRVKVTMTDMFLNLAIADLLFLVLPFWATAAADQWKFQTFMCKVNSMYKMFYS 132
Qy 121 CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKMVCFTIWLAAALCIPILYSQIKEE 180
|||||
133 CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKMVCFTIWLAAALCIPILYSQISGE 192
Qy 181 SGIAICTMVPSPDESTKLKSAVLTKVILGFPLPVVMACCYTIITHTLQAKSSKHKA 240
|||||
193 SGIAICTMVPSPDKNAKLKSAVLTKVILGFPLPVVMACCYTIITHTLQAKSSKHKA 252
Qy 241 LKVTITVTLVFLVSQFPYNCILLVOTIDAYAMFISNCAVSNIDICFQVQTIAFFHSCL 300
|||||
253 LKVTITVTLVFLVSQFPYNCILLVOTIDAYAMFISNCTISTNIDICFQVQTIAFFHSCL 312
Qy 301 NPVLVYVGERFRRLDLVTKNLGICISQAQWVSFTRREGSKLSSMLLETSGALS 357
|||||
313 NPVLVYVGERFRRLDLVTKNLGICISQAQWVSFTRREGSKLSSMLLETSGALS 369
Db

RESULT 3
CKR7_HUMAN STANDARD; PRT; 378 AA.
ID CKR7_HUMAN

```

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 1, 2002, 06:28:56 ; Search time 28.28 Seconds
(without alignments)
2183.847 Million cell updates/sec

Title: US-09-522-752-2
Perfect score: 1854
Sequence: 1 MADDYGSSTSMEDYVNFN.....EGSLKLSMILETTSGALS 357

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 562222 seqs, 172994929 residues

Total number of hits satisfying chosen parameters: 562222

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

- 1: sp_archaea.*
- 2: sp_bacteria.*
- 3: sp_fungi.*
- 4: sp_human.*
- 5: sp_invertebrate.*
- 6: sp_mammal.*
- 7: sp_mhc.*
- 8: sp_organelle.*
- 9: sp_phase.*
- 10: sp_plant.*
- 11: sp_rodent.*
- 12: sp_virus.*
- 13: sp_vertebrate.*
- 14: sp_unclassified.*
- 15: sp_virus.*
- 16: sp_bacteriaph.*
- 17: sp_archaeap.*

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1854	100.0	369	4 Q9UQ06	Q9uqg6 homo sapien
2	776.5	41.9	368	13 O42444	O42444 oncorhynchu
3	671.5	36.2	351	11 Q9EQ16	Q9eq16 mus musculus
4	665.5	35.9	351	11 Q9ERH5	Q9erh5 mus musculus
5	645	34.8	350	11 Q924I3	Q924i3 mus musculus
6	621	33.5	343	6 Q9N020	Q9n020 cercocebus
7	616	33.2	342	6 Q9TV16	Q9tv16 pan troglod
8	616	33.2	352	6 Q9TV44	Q9tv44 cercopithec
9	614	33.1	367	11 Q9RLV0	Q9rlv0 mus musculus
10	613	33.1	343	6 Q9BDS6	Q9bds6 macaca fasc
11	612	33.0	342	4 Q9HCAS	Q9hcas homo sapien
12	612	33.0	352	6 Q9XT76	Q9xt76 cercopithec
13	612	33.0	352	6 Q95ND1	Q95nd1 mandrillus
14	611	33.0	352	6 Q9BGN6	Q9bgn6 cercopithec
15	608	32.8	352	6 Q95ND2	Q95nd2 mandrillus
16	607	32.7	352	6 Q9TV49	Q9tv49 cercocebus

ALIGNMENTS

RESULT 1

Q9UQ06 ID Q9UQ06 PRELIMINARY; PRT; 369 AA.
AC Q9UQ06
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-OCT-2001 (TREMBLrel. 18, Last annotation update)
DE CHEMOKINE RECEPTOR CCR9 (CC CHEMOKINE RECEPTOR 9A).
GN CCR9.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP MEDLINE-99248139; PubMed-10229797;
RX Zaballos A., Gutierrez J., Varona R., Ardavin C., Marquez G.;
RT "Identification of the orphan chemokine receptor GPR-9-6 as CCR9, the
RT receptor for the chemokine TECK";
RL J. Immunol. 162:5671-5675(1999).
RN [2]
RP SEQUENCE FROM N.A.
RA Yu C.-R., Peden K.W.C., Farber J.M.;
RT "CCR9A and CCR9B, Two Receptors for the Chemokine CCL25 (TECK/Ckbeta-
RT 15).";
RL Submitted (APR-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ132337; CAB3477.1; -
DR EMBL; AF145439; AAF66699.1; -
DR InterPro; IPR004069; Chemokine9_receptor.
DR InterPro; IPR000276; GPCR_Rhodpsn.
DR Pfam; PF00001; 7tm.1; 1.
DR PRINTS; PR01531; CHEMOKINER9.
DR PRINTS; PR02237; GPCR_RHODPSN.
DR PROSITE; PS00237; G_PROTEIN_RECEPTOR.
DR PROSITE; PS0262; G_PROTEIN_RECEPTOR.
KW Receptor.
SQ SEQUENCE 369 AA; 42015 MW; F27CEA0CFB6B44C CRC64;

09tuq6 erythrocebu
095ne1 cercocebus
077776 cercocebus
09tuq7 cercopithec
09tsn2 macaca fasc
097975 macaca arct
09tsk1 cercopithec
09tv42 cercopithec
09xs99 gorilla gor
095ne8 cercopithec
095nd0 erythrocebu
09xt12 cercopithec
09bgn5 cercopithec
09tv50 pan troglod
09tuw4 pan troglod
09tus6 papio cynoc
09tus5 papio cynoc
09tur4 mandrillus
095nc1 theropithec
09txq3 mandrillus
09txq2 erythrocebu
09tqus cercopithec
09tq4 cercopithec
09tq2 gorilla gor
09tuw5 pan troglod
09tut6 macaca neme
09tqx0 cercopithec
095nc5 hylobates s
095nc0 hylobates m

